In the Claims:

Please amend claims 6, 9, 10, 12-17, 19-22, 28-34, and 36-38.

Please cancel claims 1-5, 7, 8, 11, 23-27, and 35.

Please add new claims 39-45.

- 1. (Canceled)
- 2. (Canceled)
- 3. (Canceled)
- 4. (Canceled)
- 5. (Canceled)
- 6. (Currently amended) The recombinant ehimeric inhibitor protein of a protease of claim 39 5, wherein eharacterized in that the kallikrein is an hK2 kallikrein protein.
- 7. (Canceled)
- 8. (Canceled)
- 9. (Currently amended) The recombinant ehimerie inhibitor protein of a protease of claim 39 8, wherein characterized in that the serpin sequence is selected from the group consisting of comprising the α-lantichymotrypsin (ACT), protein C inhibitor (PCI), α-lantiproteinase (AAT), human α-lantitrypsin-related protein precursor (ATR), α-2-plasmin inhibitor (AAP), human anti-thrombin-III precursor (ATIII), protease inhibitor 10 (PI10), human collagen-binding protein 2 precursor (CBP2), protease inhibitor 7 (PI7), protease inhibitor leuserpin 2 (HLS2), human plasma protease C1 inhibitor (C1 INH), monocyte/neutrophil elastase inhibitor (M/NEI), plasminogen activator inhibitor-3 (PAI3), protease inhibitor-1 precursor endothelial (PAI-1), human plasminogen activator inhibitor-2 placental (PAI2), human pigment epithelium-derived factor precursor (PEDF), protease inhibitor 6 (PI6), protease

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inhibitor 8 (PI8), protease inhibitor 9 (PI9), human squamous cell carcinoma antigen 1 (SCCA-1), human squamous cell carcinoma antigen 2 (SCCA-2), T4-binding globulin (TBG), Megsin, and protease inhibitor 14 (PI14), fragments thereof, molecular chimeras thereof, combinations thereof, and/or variants thereof.

10. (Currently amended) The recombinant chimeric inhibitor protein of claim 39 protease of any of the preceding claims, wherein characterized in said recombinant chimeric inhibitor protein of a protease is selected from the group consisting of comprising MD820, MD 62, MD 61, MD 67, and MDCI.

11. (Canceled)

- 12. (Currently amended) An purified and isolated DNA sequence encoding the recombinant ehimeric inhibitor protein of claim 39 of a protease according to any of the preceding claims.
- 13. (Currently amended) The purified and isolated DNA sequence of claim 12, wherein said characterized in that the sequence is selected from the group consisting of comprising SEQ ID N° 1, SEQ ID N° 3, SEQ ID N° 5, SEQ ID N° 7, SEQ ID N° 9, SEQ ID N° 11, and SEQ ID N° 13.
- An expression vector comprising characterized in that it 14. (Currently amended) comprises the purified and the isolated DNA sequence of claims 12 to 13.
- 15. (Currently amended) The expression vector of claim 14, characterized in that it further comprising comprises a promoter operably linked to the purified and isolated DNA sequence.
- 16. (Currently amended) A eukaryotic or prokaryotic host cell transfected with the expression vector of claim elaims 14 or 15.

17. (Currently amended) A pharmaceutical composition comprising the recombinant eharacterized in that it comprises a chimeric inhibitor protein of a protease of claim 39 any of claims 1 to 11 as an active agent, and a optionally in combination with one or more pharmaceutically acceptable carriers.

- 18. **(Original)** A method of treating or preventing a proteolysis-associated disorder in a mammal comprising administering to said mammal the pharmaceutical composition of claim 17.
- 19. (Currently amended) The method of claim 18, wherein characterized in that the disorder is a disorder in which hK2 kallikrein activity is detrimental.
- 20. (Currently amended) The method of claim 18 or 19, wherein characterized in that the disorder is selected from the group consisting of a cancer, an autoimmune disorder, an inflammatory disorder, and or an infectious disorder.
- 21. (Currently amended) The method of claim 20, wherein characterized in that the cancer is selected from the group consisting of prostate cancer, breast cancer, and or a metastasic cancer.
- 22. (Currently amended) The method of claim 20, wherein characterized in that the inflammatory disorder is Benign Prostatic Hypertrophy.
- 23. (Canceled)
- 24. (Canceled)
- 25. (Canceled)
- 26. (Canceled)
- 27. (Canceled)

28. (Currently amended) A method for producing the <u>recombinant ehimerie</u> inhibitor protein <u>of claim 39</u> of a protease of claims 1-to 11, comprising the steps of

- a) selecting a polynucleotidic sequence encoding a substrate <u>active</u>-enzyme interaction site specific for said Kallikrein a protease;
- b) introducing said polynucleotidic sequence into a sequence encoding <u>a serpin</u>, so as to <u>obtain a recombinant inhibitor protein an inhibitor protein of a serine or cysteine</u> protease, so as to obtain a chimeric sequence;
- c) allowing expression of said <u>recombinant inhibitor protein</u> ehimeric sequence in a cell expression system under suitable conditions; and
- d) and recovering said recombinant the chimeric inhibitor protein of a protease.
- 29. (Currently amended) The method of claim 28, wherein characterized in that step a) is performed by phage-displayed library screening.
- 30. (Currently amended) The method of claims 28 and 29, characterized in that wherein the suitable conditions comprise consist in culturing the cell expression system at a temperature between 10-40°C during 10-30 hours.
- 31. (Currently amended) The method of claim 30, characterized in that wherein the suitable conditions comprise consist in a temperature of 16°C during 16 hours.
- 32. (Currently amended) The method of claims 28 to 31, wherein step d) characterized in that step b) is achieved by separation after extraction of said chimeric recombinant inhibitor protein of a protease from the cell expression system.
- 33. (Currently amended) The method of claim 32, wherein characterized in that the separation of said chimeric recombinant inhibitor protein of a protease is achieved by affinity chromatography.
- 34. (Currently amended) The method of claims 28 to 33, wherein characterized in that the chimeric recombinant inhibitor protein of a protease is further assayed for its ability to inhibit the activity of said kallikrein the protease.

35. (Canceled)

36. (Currently amended) The method of claim 28 35, wherein the cell expression system characterized in that the prokaryotic cell is a bacterial cell.

- 37. (Currently amended) A diagnostic kit for the detection of a <u>kallikrein protease</u> in a specimen <u>comprising</u> characterized in that it comprises any suitable purified and isolated a DNA sequence selected from the group <u>consisting of comprising</u> SEQ ID N° 1, 3, 5, 7, 9, 11, 13, a sequence complementary thereof, fragments thereof, and/or variants thereof.
- 38. (Currently amended) A diagnostic kit for the detection of a <u>kallikrein protease</u> in a specimen <u>comprising characterized in that it comprises a the recombinant chimeric inhibitor protein of claim 39 a protease according to claims 1 to 11.</u>
- 39. (New) A recombinant inhibitor protein of a kallikrein comprising a serpin sequence wherein, the Reactive Serpin Loop (RSL) of said serpin sequence is modified by at least one substrate active site sequence, fragments thereof, a molecular chimera thereof, a combination thereof, and variants thereof, specific for said kallikrein.
- 40. (New) A recombinant inhibitor protein of a kallikrein hK2, comprising a serpin sequence, wherein the Reactive Serpin Loop (RSL) of said serpin sequence is modified by at least one substrate active site sequence, fragments thereof, a molecular chimera thereof, a combination thereof, and variants thereof, specific for said kallikrein hK2.
- 41. **(New)** A recombinant inhibitor protein of a kallikrein, comprising a serpin sequence, wherein the amino acid sequence of the Reactive Serpin Loop (RSL) of said serpin sequence is selected from the group consisting of SEQ ID No 16, 17, 18, 19, 20, 21, and 22, fragments thereof, molecular chimeras thereof, combinations thereof, and variants thereof.
- 42. **(New)** The recombinant inhibitor protein of claim 39, wherein the at least one substrate active site sequence specific for said kallikrein is a substrate peptide selected by said kallikrein using a phage-displayed random pentapeptide library.

43. (New) A chimeric inhibitor protein of a kallikrein comprising a serpin sequence wherein, the Reactive Serpin Loop (RSL) of said serpin sequence is modified by at least one substrate active site sequence, fragments thereof, a molecular chimera thereof, a combination thereof, and variants thereof, specific for said kallikrein.

- 44. **(New)** A purified and isolated DNA sequence encoding the recombinant inhibitor protein of claim 39.
- 45. (New) A purified and isolated DNA sequence encoding the chimeric inhibitor protein of claim 43.